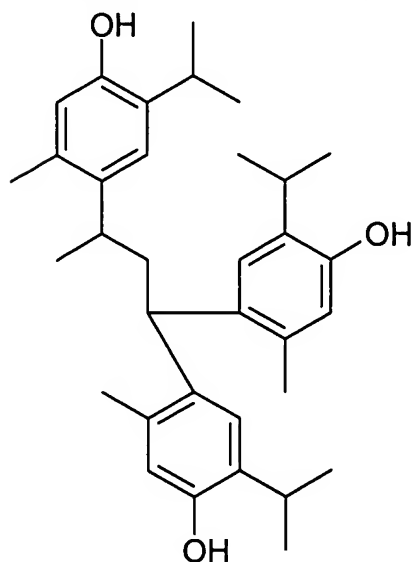


In the Claims

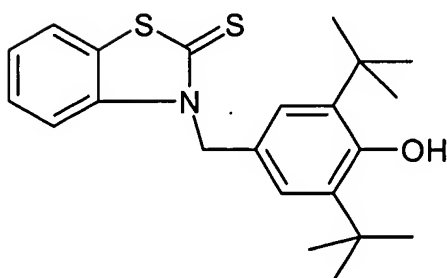
1. **(original)** Method of coloring a polymeric material, wherein a polymeric material containing

- a) a phenolic antioxidant and/or a phenolic ultraviolet absorber and
- b) a colour former

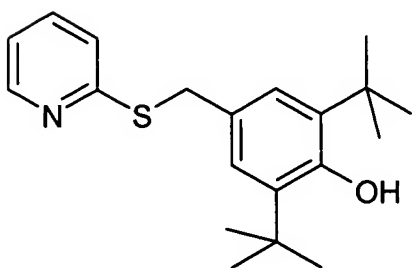
is irradiated using a radiation of higher energy than visible light, provided that the phenolic antioxidant and/or phenolic ultraviolet absorber (a) is not a compound of the formula (2) to (14)



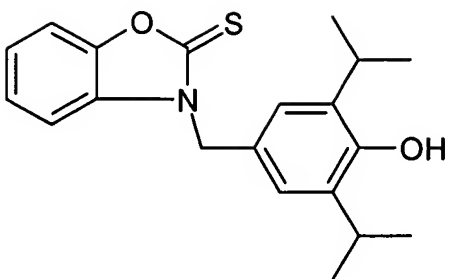
(2)



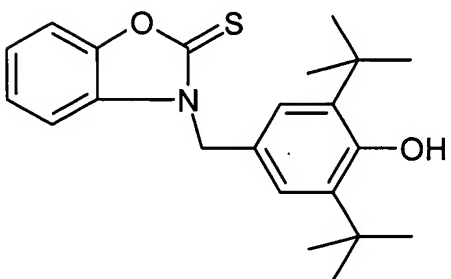
(3)



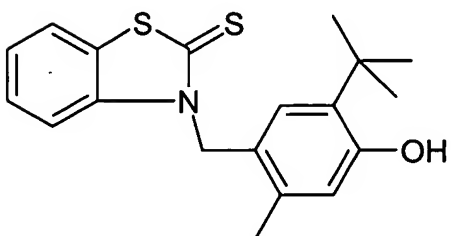
(4)



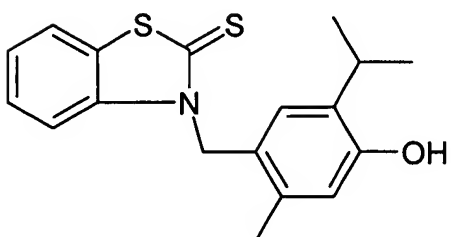
(5)



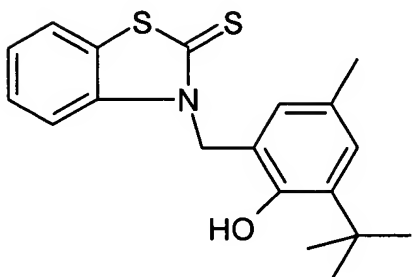
(6)



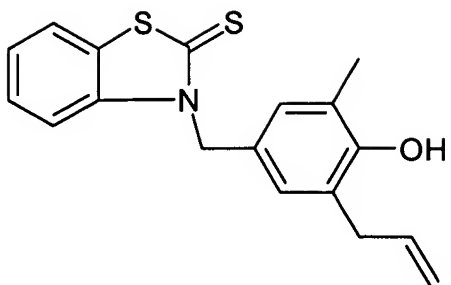
(7)



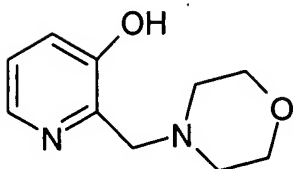
(8)



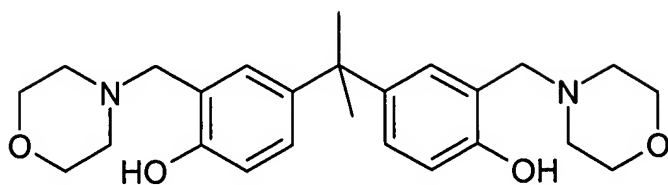
(9)



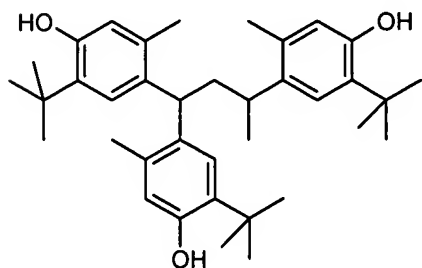
(10)



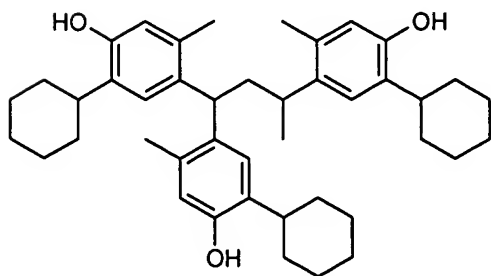
(11)



(12)



(13)



(14).

2. (currently amended) Method according to claim 1, wherein the radiation of higher energy than visible light is selected from ultraviolet light, X-ray, gamma radiation and particle radiation[**[,]**]

~~especially from ultraviolet laser or ultraviolet lamp radiation of 285 to 400 nm, electron radiation, X-ray and gamma radiation.~~

3. (currently amended) Method according to claim 1, wherein component (a) is a compound comprising one or more mono-hydroxyphenyl moieties, each carrying one or two bonds to either a linking group connecting the moiety with 1 to 3 further moieties of the same type or to an anchor group,

and optionally 1-3 further substituents selected from alkyl of 1 to 12 carbon atoms,

where the linking groups are di-, tri- or tetravalent aliphatic groups of 1 to 20 carbon atoms and

divalent linking groups are selected from

alkylene which may be interrupted and/or end-capped with -O-, -NH-, -S-, -CO-, -COO-, -OCO-,

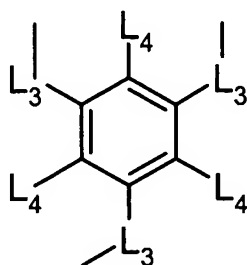
-NHCO-, -CONH-, a group L₁, phenylene~~[[.]]~~ or phenylene which is substituted by C₁-C₁₂alkyl and/or C₁-C₁₂alkoxy and/or C₂-C₁₂alkanoyloxy and/or C₃-C₁₂alkenoyloxy;

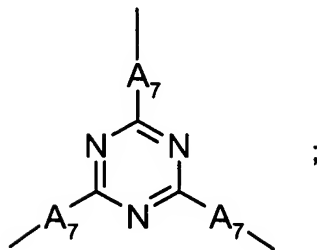
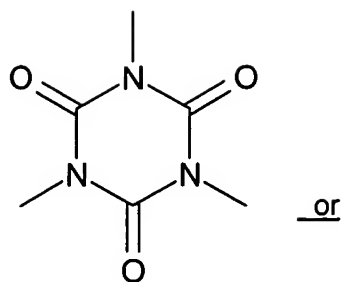
divalent mono-, di- or tricycloalkylene groups; divalent mono-, di- or tricycloalkylene groups interrupted by -O-; ~~spacer groups~~ -O-, -NH-, -S-, -CO-, -COO-, -OCO-, -NHCO-; and -CONH-;

trivalent groups are selected from

trivalent alkyl groups of 3 to 20 carbon atoms; said trivalent alkyl groups interrupted and/or end-

capped with -O-, -NH-, -S-, -CO-, -COO-, -OCO-, -NHCO-, -CONH-, a group L₁, phenylene~~[[.]]~~ or phenylene which is substituted by C₁-C₁₂alkyl and/or C₁-C₁₂alkoxy and/or C₂-C₁₂alkanoyloxy and/or C₃-C₁₂alkenoyloxy; and/or trivalent groups of the formulae



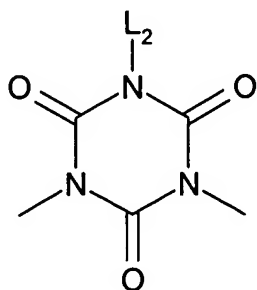


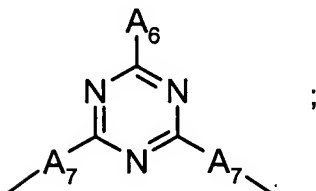
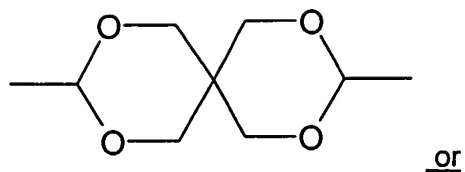
tetravalent groups are selected from

tetravalent alkyl groups of 4 to 20 carbon atoms; and said tetravalent alkyl groups interrupted and/or end-capped with -O-, -NH-, -S-, -CO-, -COO-, -OCO-, -NHCO-, -CONH-, a group L₁, phenylene[[,]] or phenylene which is substituted by C₁-C₁₂alkyl and/or C₁-C₁₂alkoxy and/or C₂-C₁₂alkanoyloxy and/or C₃-C₁₂alkenoyloxy;

wherein

L₁ is a group selected from the formulae





L₂ is OH, C₁-C₁₂alkyl, C₁-C₁₂alkoxy, C₂-C₁₂hydroxyalkyl; or C₂-C₁₂hydroxyalkoxy;

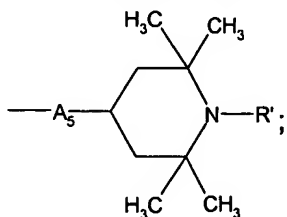
L₃ independently are C₁-C₄alkylene;

L₄ independently are H or C₁-C₄alkyl; and

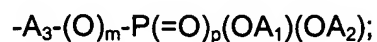
anchor groups are selected from

C₁-C₂₂alkyl; C₁-C₂₂alkyl-A₅-; C₂-C₂₂alkyl interrupted by -A₅-; -A₄-phenyl; -A₄-phenyl where the phenyl core is substituted by C₁-C₁₂alkyl, C₁-C₁₂alkoxy, C₂-C₁₂alkanoyloxy and/or C₃-C₁₂alkenoyloxy;

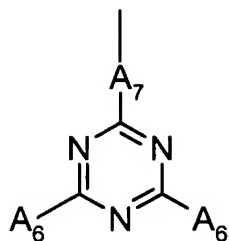
C₁-C₈alkyl substituted by a group of the formula



and phosphite, phosphate or phosphonate ester groups[[, e.g.]] of the formula



or the anchor group is of the formula



where m and p independently are 0 or 1;

A₁ and A₂ independently are C₁-C₁₂alkyl or phenyl or phenyl substituted by C₁-C₁₂alkyl or an equivalent of an alkaline, alkaline earth or aluminum atom;

A₃ is a direct bond or C₁-C₈alkylene;

A₄ is selected from C₁-C₈alkylene and A₅;

A₅ is selected from -O-, -NH-, -S-, -CO-, -COO-, -OCO-, -NHCO-**[[:]]** and -CONH-;

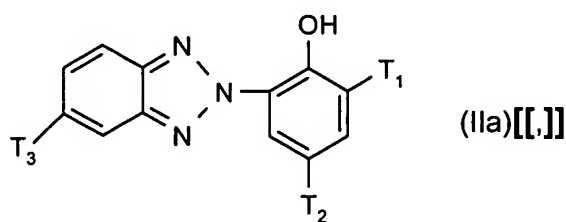
A₆ is selected from C₁-C₁₈alkoxy, C₁-C₁₈alkylthio and C₁-C₁₈alkylamino;

A₇ is -O- or -NH-;

R' is H, C₁-C₁₈alkyl, C₁-C₁₈alkoxy or cyclohexyloxy**[[:]]**;

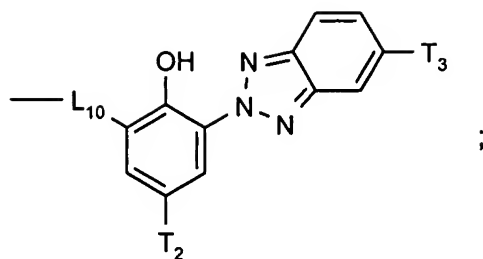
or the anchor group is C₃-C₂₂alkylene or C₃-C₂₂oxaalkylene attached with both open bonds to adjacent carbon atoms of the mono-hydroxyphenyl moiety; or

component (a) can also be a phenolic UV absorber compound selected from benzotriazoles of the formula (IIa), 2-hydroxybenzophenones of the formula (IIb)**[[:]]** and 2-hydroxyphenyltriazines of formula (IIc):



wherein T₁ is hydrogen, C₁-C₁₈alkyl, or C₁-C₁₈alkyl which is substituted by phenyl,

or T₁ is a group of the formula



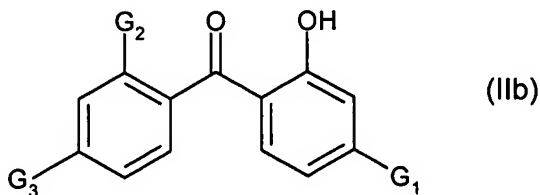
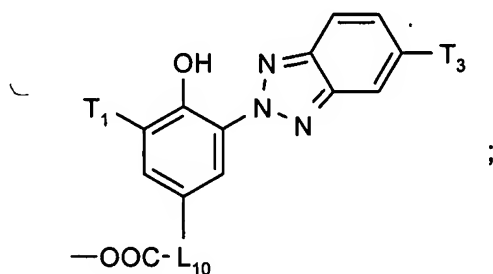
L₁₀ is a divalent group ~~[[,]]~~for example ~~-(CH₂)_n-~~, where n is from the range 1-8;

T₂ is hydrogen, C₁-C₁₈alkyl, or is C₁-C₁₈alkyl which is substituted by COOT₅, C₁-C₁₈alkoxy, hydroxyl, phenyl or C₂-C₁₈acyloxy;

T₃ is hydrogen, halogen, C₁-C₁₈alkyl, C₁-C₁₈alkoxy, C₂-C₁₈acyloxy, perfluoroalkyl of 1 to 12 carbon atoms ~~such as -CF₃,~~ or T₃ is phenyl; and

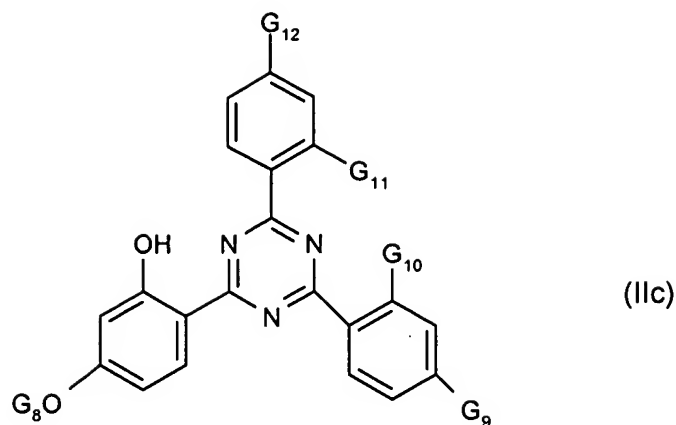
T₅ is C₁-C₁₈alkyl or C₄-C₅₀alkyl interrupted by one or more O and/or substituted by OH or by

a group



wherein

G₁, G₂ and G₃ independently are hydrogen, hydroxy or C₁-C₁₈alkoxy;

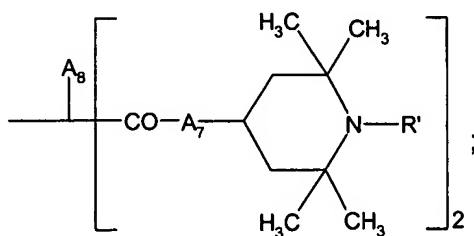


wherein

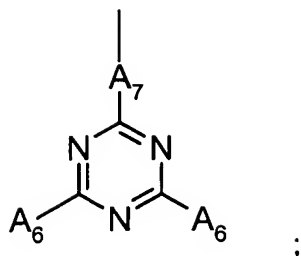
G_8 is C_1 - C_{18} alkyl, or is C_4 - C_{18} alkyl which is interrupted by COO or OCO or O, or is interrupted by O and substituted by OH; and

G_9 , G_{10} , G_{11} and G_{12} independently are hydrogen, methyl, hydroxy or OG_8 ; and G_9 and G_{12} also comprise phenyl.

4. (currently amended) Method according to claim 3, wherein the anchor groups are selected from tertiary C_4 - C_{12} alkyl; C_1 - C_{22} alkyl- A_5 -; C_2 - C_{22} alkyl interrupted by $-A_5$ -; $-A_5$ -phenyl; $-A_5$ -phenyl where the phenyl core is substituted by C_1 - C_{12} alkyl; and $-A_4$ -phenyl where the phenyl core is substituted by C_2 - C_{12} alkanoyloxy and/or C_3 - C_{12} alkenoyloxy, and optionally further by C_1 - C_{12} alkyl; or the anchor group is C_3 - C_{22} alkylene or C_3 - C_{22} oxaalkylene attached with both open bonds to adjacent carbon atoms of the mono-hydroxyphenyl moiety; or is a group of one the formulae



$-A_3-(O)_m-P(=O)_p(OA_1)(OA_2)$; or



where m and p independently are 0 or 1;

A₁ and A₂ independently are C₁-C₁₂alkyl or phenyl or phenyl substituted by C₁-C₁₂alkyl or an equivalent of an alkaline, alkaline earth or aluminum atom;

A₃ is a direct bond or C₁-C₈alkylene;

A₄ is selected from C₁-C₈alkylene, -O-, -NH-, -S-, -CO-, -COO-, -OCO-, -NHCO-**[[,]]** and -CONH-;

A₅ is selected from -O-, -NH-, -S-, -CO-, -COO-, -OCO-, -NHCO-**[[,]]** and -CONH-;

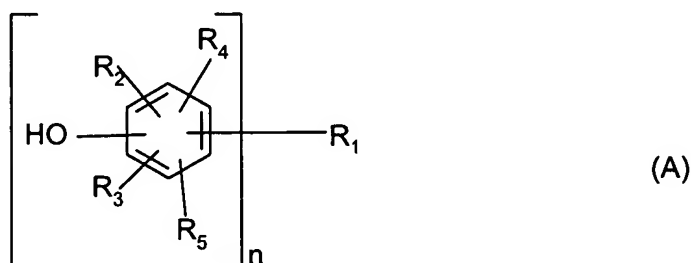
A₆ is selected from C₁-C₁₈alkoxy, C₁-C₁₈alkylthio and C₁-C₁₈alkylamino;

A₇ is -O- or -NH-;

A₈ is C₁-C₇alkyl; and

R' is C₁-C₁₈alkyl.

5. (currently amended) Method according to claim 3, wherein component (a) is a compound of the formula (A)



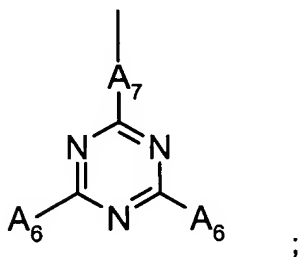
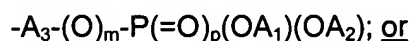
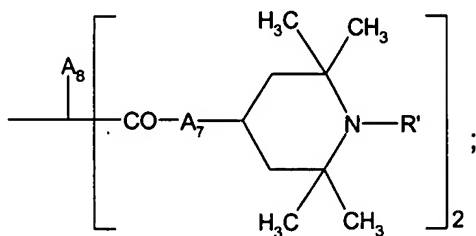
wherein

R₂, R₃, R₄ and R₅ independently are hydrogen, methyl or tertiary C₄-C₁₂alkyl**[[,]]**~~especially methyl, tert. butyl and tert. pentyl;~~

n is from the range 1-4:

when n is 1,

R₁ is tertiary C₄-C₁₂alkyl; C₁-C₂₂alkyl-A₅-; C₂-C₂₂alkyl interrupted by -A₅-; -A₅-phenyl; -A₅-phenyl where the phenyl core is substituted by C₁-C₁₂alkyl; -A₄-phenyl where the phenyl core is substituted by C₂-C₁₂alkanoyloxy and/or C₃-C₁₂alkenoyloxy, and optionally further by C₁-C₁₂alkyl; or R₁ together with R₅ is C₃-C₂₂alkylene or C₃-C₂₂oxaalkylene attached with both open bonds to adjacent carbon atoms of the mono-hydroxyphenyl moiety; or is a group of one the formulae



where m and p independently are 0 or 1;

A₁ and A₂ independently are C₁-C₁₂alkyl or phenyl or phenyl substituted by C₁-C₁₂alkyl or an equivalent of an alkaline, alkaline earth or aluminum atom;

A₃ is a direct bond or C₁-C₈alkylene;

A₄ is selected from C₁-C₈alkylene, -O-, -NH-, -S-, -CO-, -COO-, -OCO-, -NHCO-**[[.]]** and -CONH-;

A₅ is selected from -O-, -NH-, -S-, -CO-, -COO-, -OCO-, -NHCO-**[[.]]** and -CONH-;

A₆ is selected from C₁-C₁₈alkoxy, C₁-C₁₈alkylthio and C₁-C₁₈alkylamino;

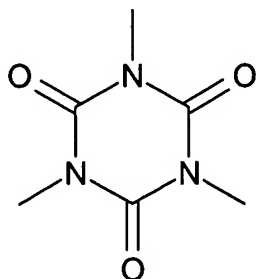
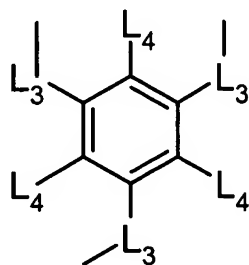
A₇ is -O- or -NH-;

A₈ is C₁-C₇alkyl; and

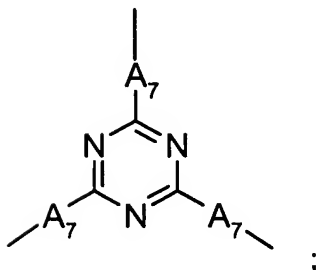
R' is C₁-C₁₈alkyl;

when n is 2, R₁ is C₁-C₂₀alkylene which may be interrupted and/or end-capped with -O-, -NH-, -S-, -CO-, -COO-, -OCO-, -NHCO-, -CONH-, -L₁-, phenylene, phenylene which is substituted by C₁-C₁₂alkyl and/or C₁-C₁₂alkoxy and/or C₂-C₁₂alkanoyloxy and/or C₃-C₁₂alkenoyloxy; divalent mono-, di- or tricycloalkylene groups; divalent mono-, di- or tricycloalkylene groups interrupted by -O-; ~~spacer groups~~ -O-; -NH-; -S-; -CO-; -COO-; -OCO-; -NHCO-; or -CONH-;

when n is 3, R₁ is trivalent alkyl of 3 to 20 carbon atoms; said trivalent alkyl interrupted or end-capped with -O-, -NH-, -S-, -CO-, -COO-, -OCO-, -NHCO-, -CONH-, -L₁-, phenylene[[,]] or phenylene which is substituted by C₁-C₁₂alkyl and/or C₁-C₁₂alkoxy and/or C₂-C₁₂alkanoyloxy and/or C₃-C₁₂alkenoyloxy; or trivalent groups of the formulae



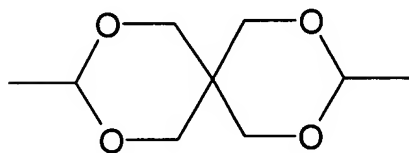
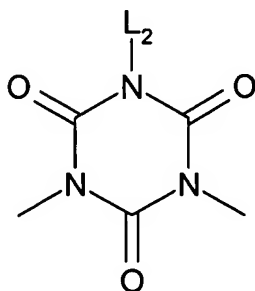
or



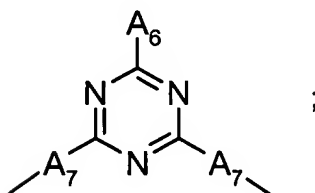
when n is 4, R₁ is tetravalent alkyl of 4 to 20 carbon atoms; said tetravalent alkyl interrupted or end-capped with -O-, -NH-, -S-, -CO-, -COO-, -OCO-, -NHCO-, -CONH-, -L₁-, phenylene[[,]] or phenylene

which is substituted by C₁-C₁₂alkyl and/or C₁-C₁₂alkoxy and/or C₂-C₁₂alkanoyloxy and/or C₃-C₁₂alkenoyloxy;

L₁ is a group selected from the formulae



and



L₂ is OH, C₁-C₁₂alkyl, C₁-C₁₂alkoxy, C₂-C₁₂hydroxyalkyl; or C₂-C₁₂hydroxyalkoxy;

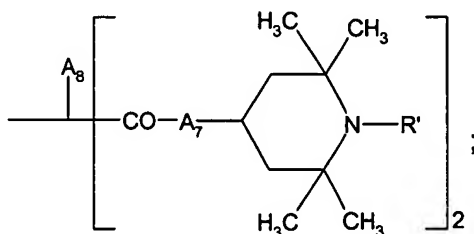
L₃ independently are C₁-C₄alkylene; and

L₄ independently are H or C₁-C₄alkyl.

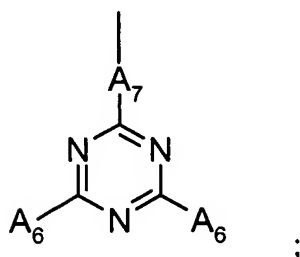
6. (currently amended) Method according to claim 5, wherein R₂, R₃, R₄ and R₅ independently are hydrogen, methyl, tert[[.]]-butyl[[.]] or tert[[.]]-pentyl;

when n is 1,

R₁ is tertiary butyl, tertiary pentyl; C₁-C₂₂alkyl-A₅-; C₂-C₂₂alkyl interrupted by -A₅-; -A₅-phenyl where the phenyl core is substituted by C₁-C₁₂alkyl; -A₄-phenyl where the phenyl core is substituted by C₃-C₄alkenoyloxy and C₁-C₁₂alkyl; or R₁ together with R₅ is C₃-C₂₂alkylene or C₃-C₂₂oxaalkylene attached with both open bonds to adjacent carbon atoms of the mono-hydroxyphenyl moiety; or R₁ is a group of one the formulae



-A₃-P(=O)(OA₁)(OA₂); or



where

A₁ and A₂ independently are C₁-C₄alkyl or an equivalent of a metal atom selected from Li, Na, K, ½ Mg, ½ Ca[[,]] and 1/3 Al;

A₃ is methylene;

A₄ is C₁-C₈alkylene;

A₅ is selected from -O-, -S-, -COO-, -OCO-, -NHCO-[[,]] and -CONH-;

A₆ is selected from C₄-C₁₈alkylthio and C₄-C₁₈alkylamino;

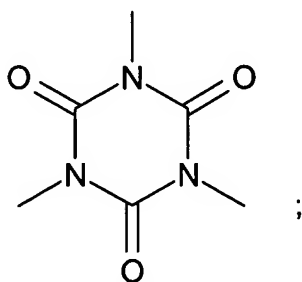
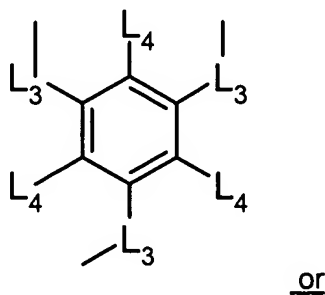
A₇ is -NH-;

A₈ is C₁-C₇alkyl; and

R' is C₁-C₁₈alkyl;

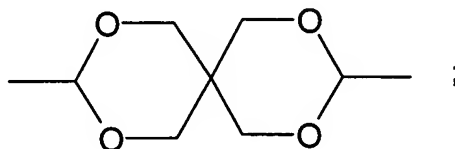
when n is 2, R₁ is C₁-C₁₂alkylene; C₂-C₂₀alkylene interrupted and/or end-capped with -O-, -S-, -COO-, -OCO-, -NHCO-, -CONH-**[[,]]** or -L₁-; or R₁ is a divalent mono-, di- or tricycloalkylene group; or R₁ is -O-; -NH-; or -S-;

when n is 3, R₁ is trivalent alkyl of 3 to 20 carbon atoms; said trivalent alkyl interrupted by -O-, -S-, -COO-, -OCO-, -NHCO-, -CONH-, phenylene**[[,]]** or phenylene which is substituted by C₁-C₁₂alkyl; or R₁ is a trivalent group of one of the formulae



when n is 4, R₁ is tetravalent alkyl of 4 to 20 carbon atoms; or said tetravalent alkyl interrupted by -O-, -S-, -COO-, -OCO-, -NHCO-**[[,]]** or -CONH-; and

L₁ is a group of the formula



L₃ independently are C₁-C₄alkylene; and

L₄ independently are H or C₁-C₄alkyl.

7. **(currently amended)** Method according to claim 1, wherein the colour former is a triphenylmethane, lactone, benzoxazine, spiropyran, ~~or preferably~~ fluoran or phthalide.

8. **(original)** Method according to claim 1, wherein the polymeric material contains 0.001 to 10 % by weight of the phenolic antioxidant and/or phenolic UVA, based on the total weight of the polymeric material.

9. **(currently amended)** Method according to claim 1, wherein the polymeric material contains 0.001 to 10 % by weight ~~[[,]] preferably 0.01 to 5 % by weight~~ of the colour former with respect to the total weight of the polymeric material.

10. **(original)** Method according to claim 1, wherein the polymeric material is a transparent thermoplast.

11. **(currently amended)** Method according to claim 1, wherein the polymeric material is selected from styrene acrylonitrile copolymer, polyolefin, polyvinylchloride, polychlorobutadiene, polyesters ~~and~~ glycol modified polyesters, polyacrylics, polystyrene, acrylonitrile styrene acrylate copolymer, polyamide, acrylonitrile styrene butadiene copolymer, polycarbonate ~~[[,]]~~ and/or blends or alloys thereof.

12. **(original)** Method of coloring a polymeric material, wherein a polymeric material containing
c) a phenolic antioxidant, phenolic ultraviolet absorber and/or a latent acid, and
d) a colour former
is irradiated using a radiation of higher energy than ultraviolet light.

13. (currently amended) Protective clothing or mask or irradiation indicating tag, wherein a polymeric material comprising components (c) and (d) according to claim 12 in the form of a fiber, textile, nonwoven or film is contained on or visibly below the surface of the clothing, mask or tag.

14. (currently amended) Process for monitoring irradiation by X-ray or radioactive material, which process comprises placing a tag or sample of a polymeric material comprising components (c) and (d) according to claim 12 in the site to be controlled, and subsequently checking the colour of the tag or sample.

15. (canceled)

16. (original) Process of making a fiber or woven or non-woven fabric, which process comprises adding (a) a phenolic antioxidant and/or phenolic UVA and (b) a colour former to a synthetic polymer before or during the fiber melt spinning process.

17. (new) Method according to claim 1, wherein the radiation of higher energy than visible light is selected from ultraviolet laser or ultraviolet lamp radiation of 285 to 400 nm, electron radiation, X-ray and gamma radiation.

18. (new) Method according to claim 1, wherein the polymeric material contains 0.01 to 5 % by weight of the colour former with respect to the total weight of the polymeric material.